

U.S. PATENT APPLICATION

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Invention: VASCULAR COATING COMPOSITION

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SPECIFICATION

VASCULAR COATING COMPOSITION

This application claims priority from Provisional Application No. 60/153,950, filed September 15, 1999, the entire content of which is incorporated herein by
5 reference.

TECHNICAL FIELD

This invention relates to methods of coating the luminal surface of a blood vessel, or other tissue cavity, and to compositions suitable for use in same.

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BACKGROUND

Various surgical and percutaneous therapies have been developed to reopen blocked channels, conduits, and other lumens, to remove diseased tissue, and to implant substitute tissue, or components thereof.

15 While these therapies are effective, they often simultaneously injure cells or cellular components of the manipulated tissues.

The responses to injuries resulting from therapeutic interventions can cause complications that
20 undo the beneficial effect of the intervention, or create new problems. For example, percutaneous transluminal angioplasty can open obstructed atherosclerotic arteries. However, the balloon-

mediated stretch and crush injury to the arterial wall can lead to proliferation of the smooth muscle cells of the media of the artery, resulting in reclosure of the artery ("restenosis") over the following months.

5 This is observed in at least one-third of arteries treated. Another example is in the formation of adhesions after surgery, in which post-operative events result in the formation or proliferation of adventitious tissue that binds internal body surfaces
10 together, causing discomfort, organ malfunction and potential morbidity and mortality. This can occur in many tissues and organs, including the intestine, the peritoneum, the heart, the pericardium, lungs, pleura, etc.

15 Various interventions have been proposed to minimize such problems. These include the use of stents and coatings in arteries (Slepian, in "Polymeric Endoluminal Paving" Cardiology Clinics 12(14) (November 1994), Slepian, et al, Circulation
20 88(4):part 2, 1-319 (1993), Hill-West, et al, Proc. Natl. Acad. Sci. USA 91:5967 (1994), USP 5,213,580, USP 5,800,538). In addition, the use of coatings, gels and fabrics can be used to prevent abdominal and pelvic adhesions (Hill-West et al, Obst. Gyne. 83:59
25 (1994)). Many of these treatments are administered after the injury has occurred, whether as a result of balloon angioplasty or otherwise, allowing the injured cells to initiate the series of processes involved in

clotting, complement activation, and cellular response
to release of cytokines, inducers of proliferation,
and other biologically active molecules. It is
difficult to stop these complex, interrelated
5 processes once they have begun.

The present invention relates to methods, and
compositions suitable for use therein, that minimize
the reaction of cells and tissues, and of cells nearby
or adjacent to them, to a subsequent injury.

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SUMMARY OF THE INVENTION

The present invention provides, in one
embodiment, a solution to the problem of restenosis
following angioplasty. Specifically, the invention
provides a method for endoluminal coating that
15 involves application of a material to the interior
surface of the involved blood vessel. The present
approach is not limited to use in connection with
restenosis, however, and can also be effectively
employed in any hollow organ to provide local
20 structural support, a smooth surface, improved flow
and sealing of lesions. In addition, the coating
material can incorporate therapeutic agents such as
drugs or cell regeneration factors to accelerate
healing processes. Such material with incorporated
25 therapeutic agents can be effectively used to coat
surgically or traumatically formed lumens in normally

solid organs as well as the native or disease generated lumens of hollow or tubular organs.

Objects and advantages of the present invention will be clear from the description that follows.

5

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1. Graph showing mean \pm sem for the BIOLASTIC disks versus the collagen-coated control dishes. Cells were counted until confluence and statistical analysis of the effect of the different growth surfaces was made. The cell present on BIOLASTIC disks were in greater number than on collagen coated plates indicating that BIOLASTIC is more suitable than collagen for endothelial cell adherence and growth.

15 Figures 2A and 2B. PKH26-GL labeled sub-confluent cultures of endothelial cells 2 days following seeding on the surface of BIOLASTIC disks (Fig. 2A) and collagen-coated plates (Fig. 2B).

20 Figures 3A and 3B. PKH26-GL labeled confluent cultures of endothelial cells 4 days following seeding on the surface of BIOLASTIC disks (Fig. 3A) and collagen-coated plates (Fig. 3B).

DETAILED DESCRIPTION OF THE INVENTION

In general, the present invention involves the introduction of a material into a selected location within a lumen in tissue, i.e. an organ, an organ
5 component or cavernous component of an organism, under conditions such that the material forms an adherent coating on the interior surface.

The basic requirements for the material to be used in the present method are biocompatibility and
10 the capacity to adhere to the surface to be coated. Suitable materials for use in the invention are described in USP 5,385,606 (Kowanko). Preferred materials comprise hemoglobin or albumin (e.g., bovine albumin) in a 30-55% solution and glutaraldehyde in a
15 5-15% solution. More preferred materials comprise bovine albumin in a 50% solution and glutaraldehyde in a 5% solution.

Coatings in accordance with the invention can be of varying thicknesses and three-dimensional
20 geometries. The coating can be present of the inner surface of hollow, cavernous, or tubular biological structures (whether natural or artificially formed) in either single or multiple layer configurations. The coating can also be used, where appropriate, to
25 occlude a tissue lumen completely. The material used can be combined with a variety of therapeutic agents for on-site delivery. Examples for use in coronary

artery applications are anti-thrombotic agents, e.g., prostacyclin and salicylates, thrombolytic agents, e.g., streptokinase, urokinase, tissue plasminogen activator (TPA) and anisoylated plasminogen-

- 5 streptokinase activator complex (APSAC), vasodilating agents, i.e., nitrates, calcium channel blocking drugs, anti-proliferative agents, i.e., colchicine and alkylating agents, intercalating agents, growth modulating factors such as interleukins,
- 10 transformation growth factor-beta and congeners of platelet derived growth factor, monoclonal antibodies directed against growth factors, anti-inflammatory agents, both steroidal and non-steroidal, and other agents that can modulate vessel tone, function,
- 15 arteriosclerosis, and the healing response to vessel or organ injury post intervention. Antibiotics can also be included in the coatings of the invention. In applications where multiple layers are used, different pharmacological agents can be used in different
- 20 layers. Moreover, the coating can be used to effect pharmaceutical delivery focally within the vessel wall.

The process of applying the coating of the invention can be carried out using catheters that may

25 incorporate occlusion balloons. The distal end resides at the delivery site and the proximal end remains outside of the patient. Coating materials can be delivered through the lumen or lumens of the

catheter. An alternative method of applying the coating is to apply a solution of, for example, glutaraldehyde using, for example, surgical gauze or sponges. The protein solution (e.g., albumin solution) can be applied as a wash onto the treated surface.

In addition to arteries, i.e., coronary, femoral, the aorta, ilial, carotid and vertebro-basilar, the present process can be utilized for other applications such as coating the interior of vascular and dialysis access graft, veins, ureters, urethrae, bronchi, biliary and pancreatic duct systems, the gut, eye, nasal passage, sinus, capsular joint, esophagus, lymphatic system, trachea and spermatic and fallopian tubes. The present method can also be used in other direct clinical applications even at the coronary level. These include but are not limited to the treatment of abrupt vessel reclosure post angioplasty, the "patching" of significant vessel dissection, the sealing of vessel wall "flaps", i.e., secondary to catheter injury or spontaneously occurring, the coating of aneurysmal coronary dilations associated with various vascular diseases. Further, the present invention provides intra-operative uses such as sealing of vessel anastomoses during coronary artery bypass grafting and the provision of a bandaged smooth surface post endarterectomy.

gently washed with Dulbecco's phosphate buffered saline (DPBS, Bio-Whittaker, Walkersville, MD) and filled with 1mg/ml type-2 collagenase in DPBS solution (216 U/mg, Worthington, Lakewood NJ) and warmed to 37°C to loosen endothelial cells from their sub-endothelial matrix. Incubation with the collagease solution for 15 minutes at 37°C was found to be adequate. The lumen was then flushed with 8-10 ml DPBS at 4°C. The eluate was centrifuged at 180 x g for 10 minutes at 4°C in tubes containing 10 ml of heat inactivated fetal bovine serum (Bio-Whittaker, Walkersville, MD) at 4°C. The supernatant was discarded and the pellet resuspended in 2 ml endothelial cell growth media (Clonetics, San Diego, CA) supplemented with 6mg bovine brain extract, 5 mg epidermal growth factor, 0.5 mg hydrocortisone, 10 ml fetal bovine serum and 25 mg gentamicin. Harvested endothelial cell numbers and viability were assessed in a 50ml aliquot of media by trypan blue exclusion. Verification of cell type was made using immunohistochemistry staining. Endothelial cells were frozen in 10% dimethylsulfoxide containing cell growth media and stored in liquid nitrogen until use.

25 *Endothelial cell maintenance culture*

Cultures of endothelial cells were maintained until use at low passage by using cryopreserved cell stocks. After thawing, frozen cell stocks were fed

three times weekly and passaged at least once a week in fresh endothelial cell growth media.

BIOLASTIC disk and collagen coating preparation

5 BIOLASTIC is comprised of crosslinked bovine albumin. To prepare BIOLASTIC disks, a 45% solution of bovine albumin was pipetted, using a pipetter mixing tip, together with a 10% solution
10 glutaraldehyde between two glass plates that were separated by 0.5 mm spacers. The ratio of albumin to glutaraldehyde present in the mixture is 4:1. This ratio initiates rapid cross-linking which is stable within 2 to 3 minutes. The 0.5 mm thick sheets were then cut into 28 mm diameter disks, which were placed
15 into sealed pouches containing water. Pouches were then sterilized using gamma irradiation (30 kGy). The disks were then removed from the sealed pouches at ambient temperature then placed in 5 ml endothelial cell growth medium for 30 minutes. Prior to seeding,
20 each disk was transferred to a 35 mm polystyrene cell culture plate (Becton Bickinson, Lincoln Park, NJ) containing 5 ml of endothelial cell growth medium.

 The control group consisted 35 mm polystyrene cell culture plates where the inside of individual
25 plates were brushed with a solution of collagen (Cohesion Technologies, Palo Alto, CA) and allowed to dry. Endothelial cells have specific substrate requirements for adherence and collagen is commonly

used as a coating material for endothelial cell culture. A 5 ml volume of endothelial cell growth medium was added prior to seeding with cells.

5 *Cell seeding and counting*

To initiate growth determinations, endothelial cells at passage 4 were carefully pipetted onto the surface of five BIOLASTIC disks and five collagen coated 35 mm control plates at a seeding density of 1.25x10⁵ cells/cm². Growth of cells attached to BIOLASTIC disks and the collagen coated control plates were monitored daily by counting cells within 4 randomly selected microscopic fields of a size 1.8 mm². Counts were facilitated by capturing video images using an inverted microscope fitted with a 4x objective and a Sony DXC-107 digital camera and a UP5000 video printer. Printed field images were manually counted and an average of cell counts per field was calculated. From these data, growth profiles could be followed for endothelial cells seeded onto BIOLASTic disks and the control collagen-treated culture plates. All results were calculated as the mean number of cells and the standard error from the mean. Statistical differences between groups were calculated using the Student's t test.

Endothelial cell staining

At various times during the seeding experiments cultures of endothelial cells growing on BIOLASTIC disks and collagen coated plates were stained with a fluorescent lipophilic dye to visualize the cell morphology. Rapid *in situ* staining of endothelial cells is now available using the fluorescent lipophilic compound, PKH-26-GL. With minimal sample manipulation the endothelial cell coverage is clearly seen under fluorescent light. BIOLASTIC disks were removed from endothelial cell culture medium and labeled with the PKH26-GL dye (Sigma Chemical Company, St. Louis, MO) according to Horan et al (Methods Cell. Biol. 3:469-490 (1990)). Collagen coated plates were stained in a similar way. Briefly, the disks or plates were first rinsed with DPBS prior to labeling with 200 ml of 10 μ M PKH26-GL. After a 10 minute incubation, the disks and plates were rinsed in fetal bovine serum and then with DPBS. The microscope used for visualization was fitted with an epi-fluorescent attachment and a Nikon G-series filter set.

Results

Endothelial cell staining

Staining of endothelial cell cultures with the lipophilic dye PKH26-GL was rapid. Cells were visualized within 15 minutes of initiating the staining protocol and showed clear cell outlines,

cell-cell contact points and cell nuclei (Fig. 2, Fig. 3).

Growth of cell cultures

5 After seeding, cells on the BIOLASTIC disks showed good cell attachment to the surface within six hours. This was also seen in collagen-coated plates. On the day following seeding, the cells flattened on the surface of both the BIOLASTIC disks and the
10 collagen-coated plates. They exhibited a characteristic polygonal and elongated endothelial cell morphology (Fig. 2A, Fig. 2B). A rapid increase in cell number was seen in the first two days following seeding of the BIOLASTIC disks and the
15 collagen-coated plates (Table 1, Fig. 1). Between days 2 and 4 after seeding, endothelial cells continued to proliferate at similar rates on both the BIOLASTIC disks and the collagen-coated plates (Table 1, Fig. 1). After day 4, cell populations attained
20 confluence and complete coverage of the BIOLASTIC disks and the collagen coated plates by a monolayer of cells (Fig. 3A, Fig. 3B). Beyond this time, no further increase in cell number was noted for either group.

25

Table 1. Growth of endothelial cells on collagen-coated plates versus BIOLASTIC disks.

| Days Following Seeding | Collagen Coated Plates n=5 | BIOLASTIC Disks n=5 |
|------------------------|-------------------------------|------------------------|
| 1 | 85 \pm 16 | 161 \pm 68 |
| 2 | 129 \pm 54 | 204 \pm 64 |
| 3 | 185 \pm 70 | 286 \pm 76 |
| 4 | 230 \pm 85 | 351 \pm 98 |

5

All values are reported as the mean \pm standard error from the mean.

Statistics

From the time of initial cell seeding to
 10 confluence of cell cultures, there was a greater
 number of cells on BIOLASTIC disks than the collagen-
 coated control plates. With five pairs of data at
 each time point, this difference attained statistical
 significance at the 1% level at all four time points
 15 ($p < 0.01$) as performed by Student's two sample unpaired
 t-test.

When the combined data for the four days
 following seeding was taken together and analyzed
 using analysis of variance, the difference between the

number of cells present on the BIOLASTIC disks compared to the collagen-coated plates was highly significant, $p < 0.001$, with 7 degrees of freedom.

5

EXAMPLE 2

Experimental details

A 50% solution of bovine albumin was made by dissolving 25g bovine albumin in 50ml deionized water at 37°C to facilitate solubilization of the albumin.

10 A 5% solution of glutaraldehyde was prepared from a stock glutaraldehyde solution of 25%. To create a thin coating on the luminal surface of a small diameter vascular graft, 10 ml of glutaraldehyde pre-warmed to 37°C was run, over a period of 30 seconds,

15 down the lumen of a fresh pig carotid artery. This was achieved by holding one end of the artery by a cannula inserted and tied in place for support. Immediately following glutaraldehyde, 10ml of albumin solution, pre-warmed to 37°C, was run down the lumen

20 of the carotid artery over 30 seconds in a similar way to the glutaraldehyde solution. Immediately following the albumin, 10ml of phosphate buffered saline at 37°C was run through the lumen of the carotid artery over a 30 second period as a wash solution. Cross-sections

25 of the carotid artery were then cut for examination of the surface coating integrity at various places along

the length of the vessel. Similar coating treatments were carried out in fresh porcine and bovine ureters.

Results

5 The thickness of the glutaraldehyde/albumin coating was measured using a Nikon microscope fitted with a digital camera. Video images were measured using a computerized morphometric software routine to yield an average thickness of the coating. In general, coating thickness measurements yielded an
10 average coating thickness of 28 microns (0.028 mm).

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All documents cited above are hereby incorporated in their entirety by reference.

15 One skilled in the art will appreciate from a reading of this disclosure that various changes in form and detail can be made without departing from the true scope of the invention.